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NEWS	4	FEB	02	GENBANK enhanced with SET PLURALS and SET SPELLING
NEWS	5	FEB		Patent sequence location (PSL) data added to USGENE
NEWS		FEB		COMPENDEX reloaded and enhanced
NEWS		FEB		WTEXTILES reloaded and enhanced
NEWS	8	FEB	19	New patent-examiner citations in 300,000 CA/CAplus patent records provide insights into related prior art
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NEWS	10	FEB	23	Several formats for image display and print options discontinued in USPATFULL and USPAT2
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NEWS	12	FEB	23	TOXCENTER updates mirror those of MEDLINE - more precise author group fields and 2009 MeSH terms
NEWS	13	FEB	23	Three million new patent records blast AEROSPACE into STN patent clusters
NEWS	14	FEB	25	USGENE enhanced with patent family and legal status display data from INPADOCDB
NEWS	15	MAR	06	INPADOCDB and INPAFAMDB enhanced with new display formats
NEWS	16	MAR	11	EPFULL backfile enhanced with additional full-text applications and grants
NEWS	17	MAR	11	ESBIOBASE reloaded and enhanced
NEWS		MAR		CAS databases on STN enhanced with new super role
				for nanomaterial substances
NEWS	19	MAR	23	CA/CAplus enhanced with more than 250,000 patent equivalents from China
NEWS	20	MAR	30	IMSPATENTS reloaded and enhanced
NEWS	21	APR	03	CAS coverage of exemplified prophetic substances enhanced
NEWS	22	APR	07	STN is raising the limits on saved answers
NEWS	23	APR	24	CA/CAplus now has more comprehensive patent assignee
				information
NEWS	24	APR	26	USPATFULL and USPAT2 enhanced with patent assignment/reassignment information
NEWS	25	APR	28	CAS patent authority coverage expanded
NEWS		APR	28	ENCOMPLIT/ENCOMPLIT2 search fields enhanced
NEWS	27	APR	28	Limits doubled for structure searching in CAS REGISTRY

AND CURRENT DISCOVER FILE IS DATED 23 JUNE 2008.

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=> file caplus COST IN U.S. DOLLARS

COST IN U.S. DOLLARS
FULL ESTIMATED COST

SINCE FILE TOTAL ENTRY SESSION 0.22 0.22

0.50

0.72

FILE 'CAPLUS' ENTERED AT 15:27:28 ON 29 APR 2009
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FILE COVERS 1907 - 29 Apr 2009 VOL 150 ISS 18 FILE LAST UPDATED: 28 Apr 2009 (20090428/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2008.

CAS Information Use Policies apply and are available at:

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=> file reg

FULL ESTIMATED COST

COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION

FILE 'REGISTRY' ENTERED AT 15:27:32 ON 29 APR 2009 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.

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STRUCTURE FILE UPDATES: 27 APR 2009 HIGHEST RN 1140067-87-3 DICTIONARY FILE UPDATES: 27 APR 2009 HIGHEST RN 1140067-87-3

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TSCA INFORMATION NOW CURRENT THROUGH January 9, 2009.

Please note that search-term pricing does apply when conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/support/stngen/stndoc/properties.html

=> s deramciclane

L1 5 DERAMCICLANE

=> s deramciclane/cn

1 DERAMCICLANE/CN

=> d L2 str cn rn

ANSWER 1 OF 1 REGISTRY COPYRIGHT 2009 ACS on STN

Absolute stereochemistry.

- **PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**
- Ethanamine, N,N-dimethyl-2-[[(1R,2S,4R)-1,7,7-trimethyl-2phenylbicyclo[2,2,1]hept-2-vl]oxv]- (CA INDEX NAME) OTHER CA INDEX NAMES:
- CN Ethanamine, N,N-dimethyl-2-[(1,7,7-trimethyl-2-phenylbicyclo[2.2.1]hept-2yl)oxy]-, (1R-exo)-

OTHER NAMES:

CN Deramciclane 120444-71-5 REGISTRY

RN

=> file caplus COST IN U.S. DOLLARS

SINCE FILE TOTAL. ENTRY SESSION 13.23 13.95

FILL ESTIMATED COST

FILE 'CAPLUS' ENTERED AT 15:28:24 ON 29 APR 2009

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FILE COVERS 1907 - 29 Apr 2009 VOL 150 ISS 18 FILE LAST UPDATED: 28 Apr 2009 (20090428/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2008.

CAS Information Use Policies apply and are available at:

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s deramciclane L3 83 DERAMCICLANE

75 L4

=> s 120444-71-5 REG1stRY INITIATED

L5

=> s L7 and L8

Substance data SEARCH and crossover from CAS REGISTRY in progress... Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

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            75 L4
L6
            92 L3 OR L4
=> dup rem L6
PROCESSING COMPLETED FOR L6
L7
             92 DUP REM L6 (0 DUPLICATES REMOVED)
=> s sclerosis or creutzfeld
         34891 SCLEROSIS
            30 SCLEROSES
         34906 SCLEROSIS
                 (SCLEROSIS OR SCLEROSES)
            24 CREUTZFELD
1.8
         34929 SCLEROSIS OR CREUTZFELD
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92 S L7 1.9 T-10 2 L9 AND L8

=> d 1-2 ibib abs L10

L10 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2006:100738 CAPLUS

DOCUMENT NUMBER: 144:198849

TITLE: Novel dosage form comprising modified-release and immediate-release active ingredients

INVENTOR(S): Vaya, Navin; Karan, Rajesh Singh; Sadanand, Sunil; Gupta, Vinod Kumar

PATENT ASSIGNEE(S): India

SOURCE: U.S. Pat. Appl. Publ., 49 pp., Cont.-in-part of U.S.

Ser. No. 630,446. CODEN: USXXCO

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 20060024365	A1	20060202	US 2005-134633	20050519
IN 2002MU00697	A	20040529	IN 2002-MU697	20020805
IN 193042	A1	20040626		
IN 2002MU00699	A	20040529	IN 2002-MU699	20020805
IN 2003MU00080	A	20050204	IN 2003-MU80	20030122
IN 2003MU00082	A	20050204	IN 2003-MU82	20030122
US 20040096499	A1	20040520	US 2003-630446	20030729
PRIORITY APPLN. INFO.:			IN 2002-MU697	20020805
			IN 2002-MU699	20020805
			IN 2003-MU80	20030122
			IN 2003-MU82	20030122
			US 2003-630446	12 20030729

AB A dosage form comprising of a high dose, high solubility active ingredient as modified release and a low dose active ingredient as immediate release where the weight ratio of immediate release active ingredient and modified release active ingredient is from 1:10 to 1:15000 and the weight of modified release active ingredient per unit is from 500 mg to 1500 mg; a process for preparing the dosage form. Tablets containing 10 mg sodium pravastatin and 1000 mg niacin were prepared The release of sodium pravastatin after 24 h was 67.7%, and the release of miacin after 1 h was 84.1%.

L10 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2004:1156477 CAPLUS

DOCUMENT NUMBER: 142:69207

TITLE: Use of bicvclo[2.2.1]heptane derivatives for the

preparation of neuroprotective pharmaceutical

compositions Gacsalyi, Istvan; Gigler, Gabor; Harsin, Laszlo Gabor;

Levay, Gyoergy; Moricz, Krisztina; Simo, Annamaria;

Szenasi, Gabor

PATENT ASSIGNEE(S): Egis Gyogyszergyar Rt., Hung.

SOURCE: PCT Int. Appl., 33 pp.

CODEN: PIXXD2 DOCUMENT TYPE: Patent

LANGUAGE: English FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

INVENTOR(S):

WO 2004112769 A1 20041229 WO 2004-HU62	
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, B	
CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, E	G, ES, FI, GB, GD,
GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, K	
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, M	
NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, S	
TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, V	
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, T	
AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, C.	
EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, N	
SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, G	Q, GW, ML, MR, NE,
SN, TD, TG	
HU 2003001906 A2 20060228 HU 2003-1906	20030623
HU 2003001906 A3 20060328	
AU 2004248982 A1 20041229 AU 2004-248982	
CA 2529254 A1 20041229 CA 2004-252925	4 20040622
EP 1660063 A1 20060531 EP 2004-743721	
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, L	
IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, C	
CN 1812778 A 20060802 CN 2004-800177	
BR 2004011772 A 20060808 BR 2004-11772	20040622
ZA 2005010138 A 20070328 ZA 2005-10138 JP 2007516165 T 20070621 JP 2006-516495	20040622
MX 2005014127 A 20060224 MX 2005-14127	20040622
	20051221
NO 2006000277 A 20060119 NO 2006-277	
BG 109414 A 20061130 BG 2006-109414	
US 20060258750 A1 20061116 US 2006-562393	
PRIORITY APPLN. INFO.: HU 2003-1906	
WO 2004-HU62	
OTHER SOURCE(S): MARPAT 142:69207	20040022

$$\begin{array}{c} \text{Me} \\ \text{Me} \\ \text{O---CH}_2\text{--CH}_2\text{--N-R}^1 \\ \text{R}^3\text{---CH}_2 \end{array}$$

AB The invention relates to the use of compds. of the general Formula I, (wherein R3 stands for hydrogen or hydroxy; R1 stands for hydrogen or alkyl; and R2 stands for alkyl) and pharmaceutically acceptable acid addition salts for the preparation of pharmaceutical compns. having neuroprotective effect.

REFERENCE COUNT:

THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

>> s L6 and (AY<2004 or PY<2004 or PY<2004)
4798118 AY<2004
24035331 PY<2004
4270650 PRY<2004
L11
73 L6 AND (AY<2004 OR PY<2004 OR PRY<2004)</pre>

=> s neuroprotection or neuroprotective

9551 NEUROPROTECTION 4 NEUROPROTECTIONS 9553 NEUROPROTECTION

(NEUROPROTECTION OR NEUROPROTECTIONS)

26009 NEUROPROTECTIVE

20 NEUROPROTECTIVES 26012 NEUROPROTECTIVE

(NEUROPROTECTIVE OR NEUROPROTECTIVES)

L12 28779 NEUROPROTECTION OR NEUROPROTECTIVE

=> s L6 and L12

L13 3 L6 AND L12

=> d 1-3 L13 ibib abs

L13 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2006:100738 CAPLUS

DOCUMENT NUMBER: 144:198849

TITLE: Novel dosage form comprising modified-release and immediate-release active ingredients

INVENTOR(S): Vaya, Navin; Karan, Rajesh Singh; Sadanand, Sunil;

Gupta, Vinod Kumar

PATENT ASSIGNEE(S): India

SOURCE: U.S. Pat. Appl. Publ., 49 pp., Cont.-in-part of U.S. Ser. No. 630,446.

English

CODEN: USXXCO
DOCUMENT TYPE: Patent

LANGUAGE:

FAMILY ACC. NUM. COUNT: 2 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE		
US 20060024365	A1	20060202	US 2005-134633		20050519	
IN 2002MU00697	A	20040529	IN 2002-MU697		20020805	
IN 193042	A1	20040626				
IN 2002MU00699	A	20040529	IN 2002-MU699		20020805	
IN 2003MU00080	A	20050204	IN 2003-MU80		20030122	
IN 2003MU00082	A	20050204	IN 2003-MU82		20030122	
US 20040096499	A1	20040520	US 2003-630446		20030729	
PRIORITY APPLN. INFO.:			IN 2002-MU697	A	20020805	
			IN 2002-MU699	A	20020805	
			IN 2003-MU80	A	20030122	
			IN 2003-MU82	A	20030122	

AB A dosage form comprising of a high dose, high solubility active ingredient as modified release and a low dose active ingredient as immediate release where the weight ratio of immediate release active ingredient and modified release active ingredient is from 1:10 to 1:15000 and the weight of modified release active ingredient per unit is from 500 mg to 1500 mg; a process for preparing the dosage form. Tablets containing 10 mg sodium pravastatin and 1000 mg niacin were prepared The release of sodium pravastatin after 24 h was 67.7%, and the release of niacin after 1 h was 84.1%.

US 2003-630446

A2 20030729

L13 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2009 ACS on STN ACCESSION NUMBER: 2004:1156477 CAPLUS

DOCUMENT NUMBER: 2004:11364// CAPI

TITLE: Use of bicyclo[2.2.1]heptane derivatives for the

preparation of neuroprotective

pharmaceutical compositions

INVENTOR(S): Gacsalyi, Istvan; Gigler, Gabor; Harsin, Laszlo Gabor; Levay, Gyoergy; Moricz, Krisztina; Simo, Annamaria;

Szenasi, Gabor

PATENT ASSIGNEE(S): Egis Gyogyszergyar Rt., Hung.

SOURCE: PCT Int. Appl., 33 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.					KIND DATE			APPLICATION NO.					DATE					
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											, EC,							
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		AZ,	BY,	KG,	KZ,	MD	RU,	TJ,	TM,	AT	, BE,	BG,	CH,	CY,	CZ,	DE,	DK,	
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		SI,	SK,	TR,	BF,	BJ,	CF,	CG,	CI,	CM	, GA,	GN,	GQ,	GW,	ML,	MR,	ΝE,	
		SN,	TD,	TG														
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											, TR,							HR
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MX	2005	0141	27		A		2006	0224		MX :	2005-	1412	7		2	0051	221	
KR	2006	0239	97		A		2006	0315		KR :	2005-	7246	01		2	0051	222	
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IORIT:	Y APP	LN.	INFO	. :						HU :	2003- 2004-	1906			A 2	0030	623	
										WO :	2004-	HU62			W 2	0040	622	
HER S	OURCE	(S):			MAR	PAT	142:	6920										

$$\begin{array}{c} \text{Me} \\ \text{Me} \\ \text{Ne} \\ \text{Ne} \\ \text{Ne} \\ \text{O---} \\ \text{CH}_2 - \text{CH}_2 - \text{N--} \\ \text{R}^1 \\ \text{II} \\ \end{array}$$

AB The invention relates to the use of compds. of the general Formula I, (wherein R3 stands for hydrogen or hydroxy; R1 stands for hydrogen or alkyl; and R2 stands for alkyl) and pharmaceutically acceptable acid addition salts for the preparation of pharmaceutical compns. having neuroprotective effect.

REFERENCE COUNT:

THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2009 ACS on STN ACCESSION NUMBER: 2003:319255 CAPLUS

DOCUMENT NUMBER: 138:343854

TITLE: Buccal sprays or capsules containing drugs for treating disorders of the central nervous system

INVENTOR(S): Dugger, Harry A., III
PATENT ASSIGNEE(S): USA

SOURCE: U.S. Pat. Appl. Publ., 17 pp., Cont.-in-part of U.S. Ser. No. 537,118.

CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English

LANGUAGE: Eng FAMILY ACC. NUM. COUNT: 19 PATENT INFORMATION:

ENT	INFOR	MATI	ON:														
PA	ATENT NO.				KIND DATE			APPLICATION NO.						DATE			
	2003 9916		227		A1 A1		2003 1999			US WO	2002- 1997-	2300 US17	60 899			0020	
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		PT,	RO,	RU,	SD,						, MK, , TJ,						
	RW:	GH,		LS,	MW,						, BE,						
EP	1036	GN,					TD,	TG			2000-			cu,		9971	
	R:	ΙE,			LV,		RO				, IT,			NL,			
EP	1952				A2	D. T. F					2007-					99710	
	R:	PT,		CH,		DK,					, GR,			LI,			
EP	2042 R:			CH,	A1 DE,	DK,	2009 ES,				2008- , GR,			LI,		99710 MC,	
CA	2497				A1		2004	0429			2003-				2	0030	827
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WO	2004				A3		2004										
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											, MW,						
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	RW:										, TZ,			ZW,	AM,	AZ,	BY,
											, CH,						
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	2003		64		A1						2003-					0030	
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	2005				A1		2005				2004-					0040	
	2006				A1 A1		2006				2006-					0060: 0060:	
	2006				A1		2006				2006- 2006-					0060. 0060!	
	2006				A1		2006				2006- 2006-					0060:	
US	2006	0210	240		AI		2006	0928		US	2006-	4432	55		2	,000:	JJI

US 20060216241	A1	20060928		2006-443254		20060531
JP 2009079060	A	20090416		2008-266598		20081015
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			US	2000-537118	A2	20000329
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			JP	2000-513555	A3	19971001
			US	2002-230060	A	20020829
			WO	2003-US26847	W	20030827
			US	2003-671709	A3	20030929
			US	2003-671715	A3	20030929
			US	2003-671720	A3	20030929
			TTC	2004-034015	7.3	20040427

AB Buccal aerosol sprays or capsules using polar and non-polar solvent have now been developed which provide biol. active compds. for rapid absorption through the oral mucosa, resulting in fast onset of effect. The buccal polar compns. of the invention comprise formulation A: aqueous polar solvent, active compound, and optional flavoring agent, formulation B: aqueous polar solvent, active compound, optionally flavoring agent, and propellant; formulation C: non-polar solvent, active compound, and optional flavoring agent; and formulation D: non-polar solvent, active compound, optional flavoring agent; and propellant. Thus, a lingual spray contained sumatriptan succinate 10-15, EtOH 10-20, propylene glycol 10-15, PEG 35-40, water 10-15, and flavors 2-3%.

-4.10

-4.10

=> logoff
ALL L# QUERIES AND ANSWER SETS ARE DELETED AT LOGOFF
LOGOFF? (Y)/N/HOLD:v

CA SUBSCRIBER PRICE

COST IN U.S. DOLLARS
SINCE FILE TOTAL
ENTRY SESSION
FULL ESTIMATED COST 400.22 58.09

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL
ENTRY SESSION
ENTRY SESSION

STN INTERNATIONAL LOGOFF AT 15:38:38 ON 29 APR 2009